## **Amendments to the Claims:**

This listing of claims will replace all prior versions, and listings, of claims in the application:

## **Listing of Claims:**

Cancel claims 1-27.

- 28. (New) A pharmaceutical liposomal formulation, characterized in that it comprises as active ingredient a 3-amidino-or 3-guanidino phenylalanine derivative which is effective as urokinase inhibitor, where the active ingredient is present in a proportion by weight of 0.5-100 based on the total weight of the formulation.
- 29. (New) The formulation as claimed in claim 28, characterized in that the urokinase inhibitor is selectedfrom Na-(2,4,6-triisopropylphenyl sulfonyl)-3-amidino-(D,L)-henylalanine-4-ethoxy carbonylpiperazide,theL enantiomer thereof or a pharmaceutically suitable salt of these compounds.
- 30. (New) The formulation as claimed in claim 28, characterized in that the urokinase inhibitor is selected from Na-(2,4,6-triisopropylphenyl sulfonyl)-3-guanidino-(D,L)-phenylalanine-4 ethoxycarbonylpiperazide, the L enantiomer thereof or a pharmaceutically suitable salt of these compounds.
- 31. (New) The formulation as claimed in claim 28, characterized in that the active ingredient is present in a proportion by weight of 2-50.

- 32. (New) The formulation as claimed in claim 28, characterized in that it has a pH in the range 5.5-9.0.
- 33. (New) The formulation as claimed in claim 28, characterized in that it comprises phospholipids in a proportion by weight of 4.5-400 based on the total weight of the formulation.
- 34. (New) The formulation as claimed in claim 28, characterized in that it comprises phospholipids selected from neutral phospholipids, anionic phospholipids and combinations thereof.
- 35. (New)The formulation as claimed in claim 28, characterized in that it comprises at least one anionic phospholipids such as, for example, phosphatidylethanolamine, phosphatidylglycerol, diphosphatidylglycerol, phosphoinositol or esterified derivatives thereof.
- 36. (New) The formulation as claimed in claim 34, characterized in that it comprises phosphatidylcholine and dimyristoylphosphatidyl glycerol in a ratio of 70:30 by weight.
- 37. (New)The formulation as claimed in claim 28, characterized in that it additionally comprises a membrane-stabilizing component such as, for example,

- cholesterol, in a proportion by weight of up to 5% based on the total weight of the formulation.
- 38. (New) The formulation as claimed in claim 28 characterized in that it additionally comprises a cryoprotectant.
- 39. (New) The formulation as claimed in claim 38, characterized in that the cryoprotectant is present in a proportion by weight of up to 150, preferably 5-150, based on the total weight of the formulation.
- 40. (New) The formulation as claimed in claim 38, characterized in that the cryoprotectant is selected from carbohydrates or/and sugar alcohols.
- 41. (New) The formulation as claimed in claim 28, characterized in that the average diameter of liposomes is not greater than 500 nm.
- 42. (New) The formulation as claimed in claim 41, characterized in that the average diameter of liposomes is 100-200 nm.
- 43. (New) The formulation as claimed in claim 28, characterized in that the liposomes are unilamellar liposomes.
- 44. (New) The formulation as claimed in claim 28, for parenteral administration.

- 45. (New) The formulation as claimed in claim 44 for intravenous injection.
- 46. (New) The formulation as claimed in claim 44 for infusion.
- 47. (New) The formulation as claimed in claim 44 for subcutaneous injection.
- 48. (New) The formulation as claimed in claim 44 for subcutaneous injection
- 49. (New) The formulation as claimed in claim 28 in dehydrated form.
- 50. (New) The formulation as claimed in 28 for controlling urokinase-associated disorders.
- 51. (New) The formulation as claimed in claim 50 for controlling tumors.
- 52. (New) The formulation as claimed in claim 51 for controlling carcinomas of the breast, pancreatic carcinomas or/and the formation of metastases.
- 53. (New) The use of a formulation as claimed in claim 28 in combination with cytostatic agents.